

Culture confirmation of gonococcal infection by recall of subjects found to be positive by nucleic acid amplification tests in general practice

Jens Kjølseth Møller

► To cite this version:

Jens Kjølseth Møller. Culture confirmation of gonococcal infection by recall of subjects found to be positive by nucleic acid amplification tests in general practice. Sexually Transmitted Infections, 2010, 86 (6), pp.478. 10.1136/sti.2010.043703 . hal-00576099

HAL Id: hal-00576099 https://hal.science/hal-00576099

Submitted on 12 Mar 2011

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

1	Culture confirmation of gonococcal infection by recall of subjects found to be positive by
2	nucleic acid amplification tests in general practice
3	
4	Jens Kjølseth Møller ^{1,2}
5	
6	¹ Department of Clinical Microbiology, Lillebælt Hospital, Vejle, Denmark
7	² Department of Clinical Microbiology, Aarhus University Hospital, Skejby, Denmark
8	
9	Keywords: NAAT, Neisseria gonorrhoeae, culture, recall samples, general practice
10	
11	Revised version
12	
13	Correspondence to
14	Dr J.K. Møller, Department of Clinical Microbiology, Institute of Regional Health Services
15	Research, University of Southern Denmark, Lillebælt Hospital, Kabbeltoft 25, DK-7100
16	Vejle, Denmark; jkm@dadlnet.dk
17	

Abstract 19

gonococcal infection in the community.

20

21

22

23

Objectives: To evaluate a routine notification of general practitioners to recall nucleic acid amplification test (NAAT) positive subjects for culture of Neisseria gonorrhoeae to confirm **Methods:** A retrospective observational study of the routine testing for *N. gonorrhoeae* by

24 analysis of test results compiled from the laboratory information system in two departments 25 of clinical microbiology.

26 **Results:** Altogether, 158 male and female subjects with NAAT positive results for *N*.

27 gonorrhoeae were included in the study. Samples for culture of N. gonorrhoeae were

28 collected from 102/158 (64.6%) subjects recalled after a NAAT assay was found positive.

29 Growth of N. gonorrhoeae was seen in the samples from 54/102 (52.9%) of the re-examined

30 NAAT positive subjects. Among subjects with samples collected within the first week after

31 the positive NAAT test, 34/44 (77%) were confirmed positive by culture.

32 **Conclusion:** This study shows that it is possible for the general practitioner to recall a

33 substantial number of NAAT positive subjects to collect swabs for culture of N. gonorrhoeae

34 to confirm gonococcal infection in the community. The majority of recall samples is culture

35 positive if collected within a week of the NAAT positive test, and may provide a sufficient

36 monitoring of the drug susceptibility of N. gonorrhoeae strains in the community.

INTRODUCTION: The gold standard of laboratory diagnosis of gonorrhoea has been the
culture of the causative agent, *Neisseria gonorrhoeae* (1). An increasing number of patients in
our area are primarily diagnosed with gonorrhoea by a nucleic acid amplification test (NAAT)
only. The aim of this study was to evaluate the recall of NAAT positive subjects to collect
swabs for culture confirmation of gonococcal infection in the community.

43

44 METHODS: A retrospective observational study was performed from March 2009 to March 45 2010 including all male and female subjects diagnosed by a general practitioner (GP) with a 46 NAAT positive sample for N. gonorrhoeae (NG) only. Samples were received and examined 47 in two department of clinical microbiology from their combined catchments area of 1,000,000 people. The Gen-Probe APTIMA COMBO 2[®] (AC2) and the APTIMA[®] GC (AGC) NAAT 48 assays (Gen-Probe, San Diego, USA) were used. Positive AC2 assays were repeated and 49 50 furthermore examined with the AGC assay to confirm the initial positive result. The final 51 laboratory report routinely notified the GP that the patient should be re-examined by culture 52 to obtain an antimicrobial susceptibility pattern. 53 Swabs for culture of NG (charcoal imbibed cotton swab in a Stuart medium) were transported 54 to the laboratory on the same day or mailed by ordinary postal service, and cultured on a 55 chocolate agar plate supplemented with Polymyxin B, Lincomycin, and Amphotericin B at 56 35°C and 10% CO₂ for 48 hours. N. gonorrhoeae was identified by Gram stain and 57 conventional biochemical tests. 58 The unique Danish civil registry number (CPR) was used to cross link the laboratory results 59 of all samples for NG testing of an individual, and to establish a chronological order of the 60 samples.

62	RESULTS: A total of 158 subjects, 52 from Vejle and 106 from Aarhus, were included. All
63	initially positive AC2 assays were confirmed by repeat testing and the single analyte assay
64	AGC. Table 1 shows the distribution of female and male samples. Altogether, 102/158
65	(64.6%) NAAT positive subjects had swabs collected for culture by a recall examination. A
66	positiv culture of N. gonorrhoeae was found in 54/102 (52.9%). Twenty subjects were not
67	included in the study because they had a culture for N. gonorrhoeae collected together with
68	the positive NAAT sample, 16/20 (80%) were culture positive. Corresponding figures during
69	a two-year period at the clinic of Genito-Urinary Medicine using similar NAAT and culture
70	assays were $40/45 = 89\%$.
71	Growth of N. gonorrhoeae was seen in 34 of 44 (77%) subjects recalled within the 1st week
72	after their positive NAAT test. Corresponding figures for the 2nd week, and the 3rd week or
73	later were 13/32 (41%) and 7/21 (33%), respectively (χ^2 =15.1, degrees of freedom=2,
74	P<0.001).

75

76 **DISCUSSION:**

77 Recall samples for culture of N. gonorrhoeae were collected from two thirds of NAAT 78 positive subjects. Among subjects with samples collected within the first week after the 79 positive NAAT test, three quarters were confirmed positive by culture. The weakness of the 80 retrospective observational design of the present study is the lack of data on antibiotic treatment prior to or after the initial NAAT screening assay, which may preclude the growth 81 82 of N. gonorrhoeae in a recall culture. Furthermore that information was not available to show 83 whether failure to recall was due to subjects not attending or to GPs not routinely requesting 84 culture samples. Culture confirmation of the NAAT positive results is also challenging 85 because of the rapid loss of viability of *N. gonorrhoeae* strains during the transportation of the 86 samples from the community to the laboratory (1). Previous studies also found that about two

87	thirds of the NAAT results could be confirmed by culture (2, 3). However, urgent
88	transportation and culture of samples can increase the percentage of samples with growth of
89	N. gonorrhoeae to more than 85% (3). The present study shows that it is possible for the
90	general practitioner to recall a substantial number of NAAT positive subjects to collect swabs
91	for culture of N. gonorrhoeae to confirm gonococcal infection in the community. Provided
92	that random selections of NAAT positive subjects are re-examined by culture, recall cultures
93	may provide an accurate picture of antibiotic resistance in N. gonorrhoeae strains in the
94	community. To obtain additional follow-up samples, GPs may be requested to recall subjects
95	from whom samples were not received for culture within a week after the initial positive
96	NAAT assay.
97	In conclusion, the majority of recall samples is culture positive if collected within a week of
98	the NAAT positive test, and may provide a sufficient monitoring of the drug susceptibility of
99	N. gonorrhoeae strains in the community.
100	
101	ACKNOWLEDGEMENTS, COMPETING INTERESTS, FUNDING
102	There are no competing interests.
103	The Corresponding Author has the right to grant an exclusive licence (or non exclusive for
104	
	government employees) on a worldwide basis to the BMJ Publishing Group Ltd to
105	government employees) on a worldwide basis to the BMJ Publishing Group Ltd to permit this article (if accepted) to be published in STI and any other BMJPGL
105 106	government employees) on a worldwide basis to the BMJ Publishing Group Ltd to permit this article (if accepted) to be published in STI and any other BMJPGL products and sublicences such use and exploit all subsidiary rights, as set out
105 106 107	government employees) on a worldwide basis to the BMJ Publishing Group Ltd to permit this article (if accepted) to be published in STI and any other BMJPGL products and sublicences such use and exploit all subsidiary rights, as set out in our licence http://group.bmj.com/products/journals/instructions-for-authors/licence-forms.
105 106 107 108	government employees) on a worldwide basis to the BMJ Publishing Group Ltd to permit this article (if accepted) to be published in STI and any other BMJPGL products and sublicences such use and exploit all subsidiary rights, as set out in our licence http://group.bmj.com/products/journals/instructions-for-authors/licence-forms.
105 106 107 108 109	government employees) on a worldwide basis to the BMJ Publishing Group Ltd to permit this article (if accepted) to be published in STI and any other BMJPGL products and sublicences such use and exploit all subsidiary rights, as set out in our licence http://group.bmj.com/products/journals/instructions-for-authors/licence-forms.

111 REFERENCES

- 112 1. Iwen PC, Walker RA, Warren KL, Kelly DM, Linder J, and Hinrichs SH. 1996. Effect
- 113 of off-site transportation on detection of Neisseria gonorrhoeae in endocervical specimens.
- 114 Arch. Pathol. Lab. Med. **120:**1019-1022.
- 115 2. Gopal Rao G, Bacon L, Evans J, Dejahang Y, Hardwick R, Michalczyk P, Wong J,
- and Donaldson A. 2009. Can culture confirmation of gonococcal infection be improved in
- 117 female subjects found to be positive by nucleic acid amplification tests in community clinics?
- 118 Sex. Transm. Infect. **85:**531-533. doi: 10.1136/sti.2009.036525.
- 119 3. Rao GG, Bacon L, Evans J, Dejahang Y, Michalczyk P, Donaldson N, and Lewisham
- 120 Chlamydia and Gonoccoccus Screening Programme. 2008. Prevalence of Neisseria
- 121 gonorrhoeae infection in young subjects attending community clinics in South London. Sex.
- 122 Transm. Infect. 84:117-121. doi: 10.1136/sti.2007.026914.
- 123

125 Tables

126 Table 1. Recall of NAAT positive subjects to collect swabs for culture of *N. gonorrhoeae*

127								
127								
128		NAAT positive	Swabs for culture			Culture positive		
129		No.	No.	%	(95%CI)	No.	%	(95%CI)
130								
131	Female subjects	69	51	74	(62 - 83)	22	43	(30 - 57)
132	Urogenital swab ^a	65	49			21		
133	Urine	4	2			1		
134								
135	Male subjects	89	51	57	(47 - 67)	32	63	(49 – 75)
136	Urine	70	42			28		
137	Urethral swab	19	9			4		
138								

^a Samples from urethra, cervix and vagina